

# Structural and functional insights into the unique CBS–CP12 fusion protein family in cyanobacteria

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**Cyanobacteria are important photosynthetic organisms inhabiting a range of dynamic environments. This phylum is distinctive among photosynthetic organisms in containing genes encoding uncharacterized cystathionine  $\beta$ -synthase (CBS)–chloroplast protein (CP12) fusion proteins. These consist of two domains, each recognized as stand-alone photosynthetic regulators with different functions described in cyanobacteria (CP12) and plants (CP12 and CBSX). Here we show that CBS–CP12 fusion proteins are encoded in distinct gene neighborhoods, several unrelated to photosynthesis. Most frequently, CBS–CP12 genes are in a gene cluster with thioredoxin A (TrxA), which is prevalent in bloom-forming, marine symbiotic, and benthic mat cyanobacteria. Focusing on a CBS–CP12 from *Microcystis aeruginosa* PCC 7806 encoded in a gene cluster with TrxA, we reveal that the domain fusion led to the formation of a hexameric protein. We show that the CP12 domain is essential for hexamerization and contains an ordered, previously structurally uncharacterized N-terminal region. We provide evidence that CBS–CP12, while combining properties of both regulatory domains, behaves different from CP12 and plant CBSX. It does not form a ternary complex with phosphoribulokinase (PRK) and glyceraldehyde-3-phosphate dehydrogenase. Instead, CBS–CP12 decreases the activity of PRK in an AMP-dependent manner. We propose that the novel domain architecture and oligomeric state of CBS–CP12 expand its regulatory function beyond those of CP12 in cyanobacteria.**

crystal structure | hexamer | redox | *Microcystis aeruginosa*

Cystathionine  $\beta$ -synthase (CBS) domains are widespread structural domains, conserved either as single proteins (1, 2) or as domains in a wide range of functionally different cytosolic (3–5) and membrane proteins (6–8). The importance of CBS domains is reflected by the range of different hereditary diseases in humans caused by mutation in their sequence, including homocystinuria and retinitis pigmentosa (9, 10). Each domain folds as an antiparallel  $\beta$ -sheet flanked by helices on one side and contains the secondary structural elements  $\alpha 0$ - $\beta 1$ - $\alpha 1$ - $\beta 2$ - $\beta 3$ - $\alpha 2$  (11). They always occur in pairs, forming a so-called CBS pair or Bateman module (11, 12). All structurally characterized CBS pairs, with the exception of a single archaeal representative, assemble in homodimers containing four CBS domains, oriented in a head-to-head or head-to-tail manner (11), such as CBSX1 and CBSX2, stand-alone CBS domain proteins of plants (1, 2). CBS domains are important sensors of cellular energy status, often binding adenine nucleotides as regulatory ligands (11). Plant CBSX proteins, for instance, are important partners in the thioredoxin (Trx)-mediated redox regulation of protein activity (1). Targets of Trx-regulated processes in photosynthetic organisms include proteins related to photosynthesis and oxidative stress response (13–15). By increasing the activity of all chloroplastic Trx, plant CBSX1 enhances carbon dioxide (CO<sub>2</sub>) fixation and carbohydrate synthesis, thus affecting plant development and stabilizing cellular redox homeostasis (1).

A bioinformatic analysis of 126 cyanobacterial genomes revealed the existence of CBS domain-containing proteins C-terminally fused to the small protein CP12 (chloroplast protein) (16). While these CBS–CP12 fusion proteins are to date only found in cyanobacteria, stand-alone CP12 proteins are conserved in all oxygenic photosynthetic organisms, namely plants, algae, and cyanobacteria (17–21). CP12 is an intrinsically disordered, regulatory protein of about 80 residues. It is characterized by the AWD<sub>2</sub>VEEL core sequence (22) and two N-terminal and C-terminal redox-sensitive cysteine pairs. Under oxidizing conditions, the cysteine pairs form disulfide bridges structuring two polypeptide loops required for the inactivation of two enzymes of the CO<sub>2</sub> fixation pathway (Calvin cycle), phosphoribulokinase (PRK) and glyceraldehyde-3-phosphate dehydrogenase (GAPDH), by forming a ternary GAPDH–CP12–PRK complex (17–19). The activation of both enzymes, and consequently CO<sub>2</sub> fixation, is mediated by reduction via Trx (13), causing the dissociation of the ternary GAPDH–CP12–PRK complex. Structural information on CP12 is scarce due to its intrinsic disorder. Only about 20 residues of the C-terminal part of CP12 have been structurally characterized in a complex with GAPDH, indicating the disordered state of the N-terminal region

## Significance

Carbon fixation is arguably one of the most important metabolic processes on Earth. Stand-alone CP12 proteins are major players in the regulation of this pathway in all oxygenic photosynthetic organisms, yet their intrinsic disorder has so far hampered the capturing of a principal part of their structure. Here we provide structural insights into CP12 by investigating an uncharacterized CP12 fusion protein, CBS–CP12, which is widespread among cyanobacteria, and reveal a unique hexameric structure. Our data further extend the existing knowledge of the regulation of photosynthesis and carbon fixation by the CP12 protein family, suggesting a more versatile role of this protein family in global redox regulation, predominantly in bloom-forming cyanobacteria that pose major threats in lakes and reservoirs.

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Data deposition: The experimental structure factor amplitudes and atomic coordinates of the derived crystal structural models reported in this paper have been deposited in the Protein Data Bank, [www.pdb.org](https://www.pdb.org) (PDB ID codes 5NMU, 5NPL, and 5NVD).

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even in the presence of its binding partner under oxidizing conditions (23, 24).

In contrast to CP12, the function of cyanobacterial CBS–CP12 fusion proteins is unknown. They typically occur in addition to a stand-alone CP12 and are divided into three variants according to their number of cysteine pairs in the CP12 domain (two, one, or none) (16). The CBS pairs present in these variants share a high degree of sequence similarity (16). It was hypothesized that the fusion of CP12 and CBS domains in cyanobacteria functionally connects them, implying another layer of complexity in the regulatory roles of both CP12 and CBS domains (16).

Here we report the structural and functional characteristics of a CBS–CP12 variant from the bloom-forming cyanobacterium *Microcystis aeruginosa* PCC 7806 (hereafter *M. aeruginosa*). We show that CBS–CP12 fusion proteins are widespread in cyanobacteria and particularly associated with a thioredoxin gene in bloom-forming species. The crystal structure of CBS–CP12 includes a complete structure of the N-terminal part of CP12 and reveals that CBS–CP12 forms a hexamer, unlike any other characterized CBS domain protein. We further show coregulated light-induced gene expression of CBS–CP12 with TrxA and an AMP-dependent inhibition of PRK activity by CBS–CP12 that counteracts the regulation by TrxA.

## Results

**The CBS–CP12 Protein Family Is Widespread in Cyanobacteria.** A reassessment of the occurrence of CBS–CP12 fusion proteins in 333 cyanobacterial genomes revealed the presence of five CBS–CP12 variants, the three previously described (16) and two variants with a shorter CP12 domain, CBS–CP12-N-tr and CBS–CP12-tr (Fig. 1A). The phylogenetic survey detected 232 single CBS–CP12 proteins and demonstrated their widespread distribution in all five morphological subsections of the cyanobacterial phylum, with the exception of the entire phylogenetic subclade C1 and the model strain *Synechocystis* sp. PCC 6803 (SI Appendix, Fig. S1). Truncated CBS–CP12 variants occur in the majority of strains along with the full-length variants. As previously shown (16), all CBS–CP12 variants coexist with one or more CP12 genes.

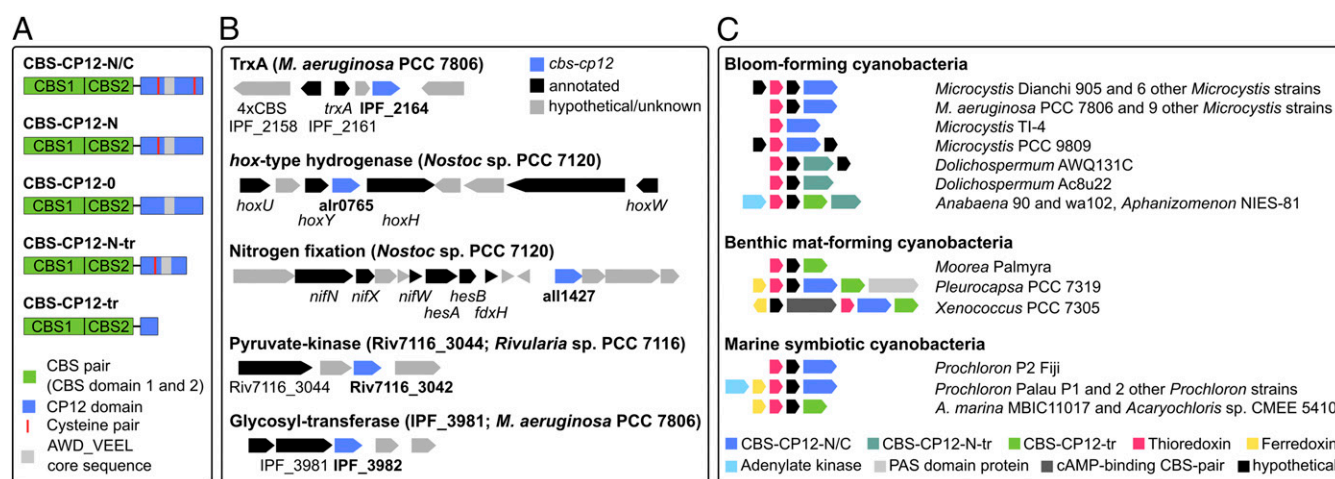
A gene neighborhood survey revealed an unexpected variety of gene clusters encoding a CBS–CP12 variant, including the *hox*-type hydrogenase and a nitrogen fixation (*nif*) gene cluster in *Nostoc* sp. PCC 7120 and a pyruvate kinase in *Rivularia* sp. PCC 7116 (Fig. 1B). Most commonly, CBS–CP12 genes were observed

in the immediate vicinity of thioredoxin A-encoding genes (Fig. 1C). These joint CBS–CP12/TrxA gene clusters were predominantly detected in genomes of bloom-forming strains, but also in benthic mat-forming and marine symbiotic cyanobacteria. Some of these gene clusters additionally encode the electron transfer protein ferredoxin and adenylate kinase, an enzyme important for cellular energy homeostasis. In all cases, a CBS protein with four CBS domains is encoded upstream of the respective gene cluster (Fig. 1B). The different CBS–CP12/TrxA gene cluster types are distributed over four cyanobacterial subclades from unicellular strains (e.g., *Microcystis* and *Acaryochloris* strains) to multicellular strains capable of forming heterocysts for nitrogen fixation (e.g., *Anabaena* and *Aphanizomenon* strains).

**CBS–CP12 Forms a Hexamer.** The CBS–CP12 protein (IPF\_2164) of the CBS–CP12/TrxA gene cluster of *M. aeruginosa* (Fig. 1) was selected as representative of the unique fusion protein family for structural analysis. It consists of 205 residues, of which about 130 N-terminal residues form a CBS pair and about 70 C-terminal residues belong to the CP12 domain, as deduced from the sequence (Fig. 2A). CBS–CP12 was expressed in *Escherichia coli* as a Trx fusion protein (SI Appendix, Table S1). Three X-ray diffraction datasets were collected: a native and an ytterbium derivative in an orthorhombic space group with three monomers in the asymmetric unit to a resolution of 2.1 and 2.8 Å, respectively, and a native in a hexagonal space group with one monomer in the asymmetric unit to 2.5 Å (SI Appendix, Table S2).

The CBS pair consists of two CBS domains, each showing a typical secondary structural arrangement (11, 12), excluding the first  $\beta$ -strand and preceded by a short  $\alpha$ -helix in the linker region (linker- $\alpha$ 1- $\beta$ 1- $\beta$ 2- $\alpha$ 2) (Fig. 2B). The  $\beta$ -strands form an antiparallel  $\beta$ -sheet, and the two  $\alpha$ -helices are located on one side of the sheet. The CBS pair is connected to the CP12 domain with a third linker of about 10 residues. Notably, the N-terminal part of CP12 is ordered (Fig. 2 and SI Appendix, Fig. S2) and forms two  $\alpha$ -helices ( $\alpha$ 5- $\alpha$ 6), of which the latter contains the AWD\_VEEL core sequence (22). Both helices are connected by an intramolecular disulfide bridge between residues Cys153 and Cys163 (Fig. 2C). About 30 C-terminal residues are not visible in the electron density.

Remarkably, the biological assembly of CBS–CP12 is a hexamer, composed of the monomers of the asymmetric unit and the symmetry equivalent (Fig. 2D and SI Appendix, Fig. S2). The



**Fig. 1.** CBS–CP12 variants and gene neighborhood conservation. (A) Domain organization of the five CBS–CP12 variants. CBS–CP12-N/C contains both cysteine pairs in the CP12 domain, CBS–CP12-N only the N-terminal pair, and CBS–CP12-0 lacks both cysteine pairs. CBS–CP12-N-tr contains a truncated CP12 domain with the N-terminal cysteine pair, and CBS–CP12-tr possesses only a short CP12 domain without cysteine pairs. The distribution of the five CBS–CP12 variants within the cyanobacterial phylum is displayed in SI Appendix, Fig. S1. (B) Selected list of gene neighborhoods of CBS–CP12 genes in different cyanobacteria. Note that a CBS–CP12 gene is also copresent in the hydrogenase gene cluster of *Synechococcus* sp. PCC 7002. (C) Conservation of 12 gene cluster types featuring a CBS–CP12 and a TrxA gene. Note that one gene cluster type occurs in bloom-forming and marine symbiotic cyanobacteria (*M. aeruginosa* PCC 7806 and *Prochloron* P2 Fiji).





CBS-CP12 (*SI Appendix, Fig. S4E*). Native MS revealed that in the presence of AMP the vast majority of CBS-CP12 appeared as a hexamer and only a small amount as a dimer, with a mass indicating the saturated binding of four ligands (*SI Appendix, Fig. S4G*).

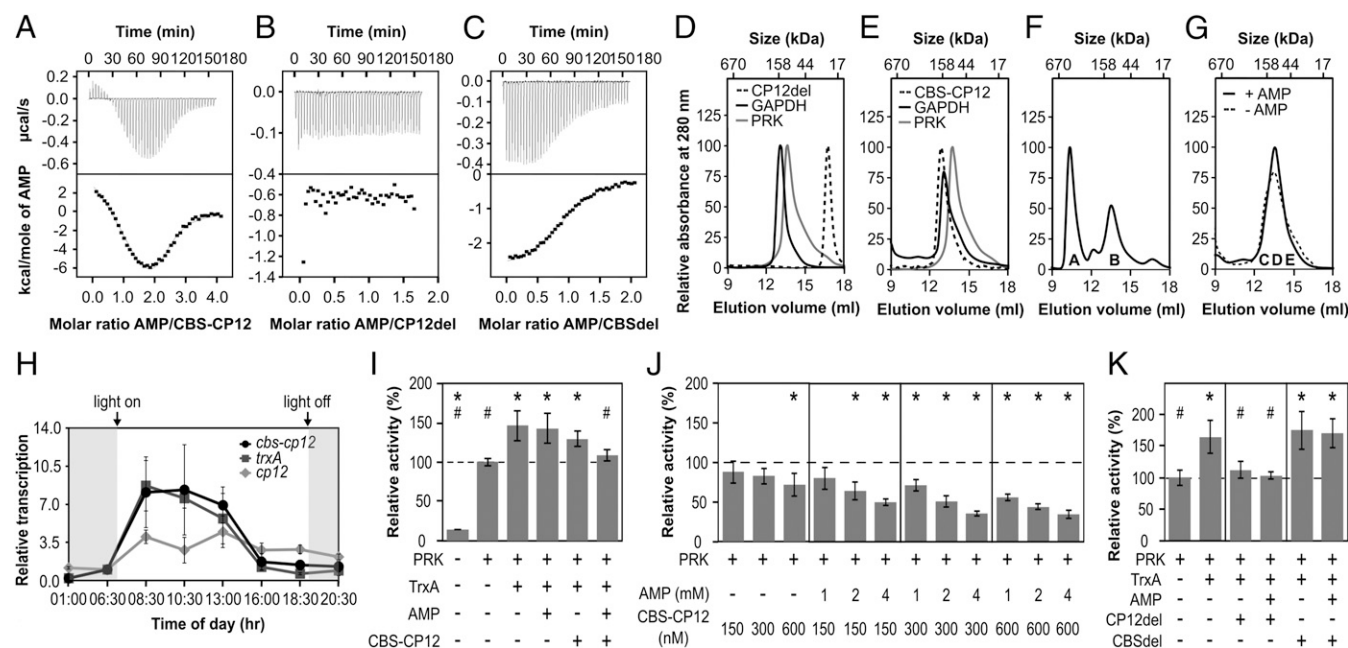
The CP12 domain CP12del did not show any interaction with AMP, while CBSdel was clearly able to bind AMP but had only one binding site (Fig. 3 *B* and *C*). CBSdel binds AMP in an exothermic reaction, similar to the reaction observed at site 2 of CBS-CP12. The binding affinity of CBSdel for AMP is about seven times lower compared with site 2, and 32 times lower for site 1 of CBS-CP12 (*SI Appendix, Table S3*). Contrary to CBS-CP12 only one AMP is bound per CBSdel monomer, indicating that two CBS domains form one AMP-binding site, as shown for CBSX2 (26).

**CBS-CP12 Does Not Replace Canonical CP12.** Using analytical SEC and CP12del as a surrogate for canonical CP12, we investigated the ability of CBS-CP12 to bind recombinant GAPDH (IPF\_4508) and PRK (IPF\_5236) of *M. aeruginosa* (*SI Appendix, Table S1*). The elution peaks of both enzymes overlap with CBS-CP12 but are separated from CP12del (Fig. 3 *D* and *E*). Under oxidizing conditions and the presence of NAD, the incubation of CP12del with GAPDH and PRK led to the emergence of a peak with an estimated mass of 569 kDa (Fig. 3*F*). Denaturing SDS/PAGE confirmed that the majority of GAPDH and PRK formed a ternary complex with CP12del (*SI Appendix, Fig. S4K*). Under the same conditions, and also with AMP, CBS-CP12 coeluted with GAPDH and PRK but did not form a ternary complex (Fig. 3*G* and *SI Appendix, Fig. S4K*). There was also no evidence from pull-down experiments for complex formation of CBS-CP12 with any other protein of *M. aeruginosa* protein extract (*SI*

*Appendix, Fig. S5 A–C*) and TrxA under different conditions (*SI Appendix, Fig. S5 D–I*).

**Cbs-cp12 and trxA Are Coexpressed and CBS-CP12 Regulates PRK Activity in Tandem with AMP.** Using RT-PCR experiments, the expression of CBS-CP12 and TrxA genes in comparison with the canonical CP12 gene (IPF\_0141) was analyzed in a 12/12-h day/night cycle (Fig. 3*H* and *SI Appendix, Table S4*). In agreement with previous data (27), canonical CP12 does not show strong variations in its gene expression. In contrast, the gene expression of CBS-CP12 and TrxA follows the same circadian rhythm and shows a clear light-induced expression, reaching their maximum early in the morning (Fig. 3*H*). Their gene expression is furthermore strongly induced under high-light stress and slightly repressed under iron-limiting conditions (*SI Appendix, Fig. S4L*).

It has been demonstrated that plant CBSX1 regulates the activity of a Calvin cycle enzyme through Trx (1). We used the Calvin cycle enzyme and Trx target PRK to examine the effect of CBS-CP12 and AMP on its activity as well as its regulation through TrxA (Fig. 3 *I–K*). TrxA increased PRK activity by 46% (Fig. 3*I*), as described previously (28). While AMP and CBS-CP12 alone did not affect this TrxA stimulation of PRK, when combined they almost completely abolished the stimulation of PRK activity by TrxA (Fig. 3*I*). This negative effect is also observed without TrxA, and depends strongly on the concentration of CBS-CP12 and AMP. Alone, AMP had only negligible effects on PRK (*SI Appendix, Fig. S4M*), while higher CBS-CP12 concentrations slightly reduced PRK activity (Fig. 3*J*). Together, however, increasing concentrations of CBS-CP12 and AMP gradually and significantly diminished PRK activity up to 65%



**Fig. 3.** Biochemical properties and gene expression of CBS-CP12. (*A–C*) ITC titration of (*A*) 400 μM AMP into 20 μM CBS-CP12, (*B*) 800 μM AMP into 100 μM CP12del, and (*C*) 700 μM AMP into 70 μM CBSdel. The black lines (*Bottom*) are the best fit to a two-site model (*A*) or one-site model (*C*). Thermodynamic data of three technical replicates are provided in *SI Appendix, Table S3*. (*D* and *E*) Individual S200 SEC elution profiles of (*D*) CP12del, GAPDH, and PRK and (*E*) CBS-CP12, PRK, and GAPDH. CP12del elutes at  $22.5 \pm 0.5$  kDa, CBS-CP12 as a hexamer at  $143.5 \pm 13.6$  kDa, PRK at  $99.2 \pm 7.5$  kDa (theoretical monomer size 38 kDa), and GAPDH at  $139.2 \pm 3.7$  kDa (theoretical monomer size 37 kDa) (mean  $\pm$  SD,  $n \geq 2$ ). (*F*) SEC elution profile of an equimolar mixture (30 μM, subunit base) of GAPDH, PRK, and CP12del with 25 mM DTTTox and 0.5 mM NAD. The ternary GAPDH-CP12-PRK complex elutes at  $\sim 569$  kDa (peak A). Peak B contains GAPDH and PRK not bound by CP12 ( $n = 1$ ). (*G*) SEC elution profile of an equimolar mixture (30 μM, subunit base) of GAPDH, PRK, and CBS-CP12 with 25 mM DTTTox, 0.5 mM NAD,  $\pm 0.5$  mM AMP. GAPDH, PRK, and CBS-CP12 coelute in fraction C-E. ( $n = 1$ ). (*F* and *G*) The denaturing SDS-PAGE of peak A and B and fractions C-E is provided in *SI Appendix, Fig. S4K*. (*H*) Diurnal rhythm of the relative transcription of *cbs-cp12*, *trxA*, and *cp12* in *M. aeruginosa* (mean  $\pm$  SD,  $n = 3$ ). The experiment was repeated twice with consistent results (*SI Appendix, Table S4*). (*I*) Relative activity of 300 nM PRK upon addition of 600 nM TrxA, 2 mM AMP, and 300 nM CBS-CP12. (*J*) Relative activity of 300 nM PRK upon addition of different concentrations of CBS-CP12 and AMP. (*K*) Relative activity of 300 nM PRK upon addition of 600 nM TrxA, 2 mM AMP, 300 nM CP12del, and 300 nM CBSdel. Data in *I–K* are mean  $\pm$  SD,  $n \geq 5$ ; \* $P \leq 0.01$  compared with 300 nM PRK, # $P \leq 0.01$  compared with 300 nM PRK + 600 nM TrxA, Student's *t* test.





GAPDH and PRK, and that CBS–CP12 fusion proteins perform additional functions. Given the widespread distribution among cyanobacteria (16), clarifying the role of CBS–CP12 proteins encoded in other gene clusters and the involvement of the other proteins encoded in the various CBS–CP12/TrxA gene cluster types (Fig. 1) will be of interest. Understanding the principles of the regulation of metabolic processes can aid in the management of harmful blooms, as caused by *M. aeruginosa*, and provide insights for metabolic engineering of photosynthesis as well as hydrogen production and nitrogen fixation in cyanobacteria.

## Materials and Methods

Cloning, expression in *E. coli*, and purification of proteins using affinity and size-exclusion chromatography, crystallization and structure determination of CBS–CP12, ITC, analytical SEC, native MS, quantitative RT-PCR, PRK and

pull-down assays, and bioinformatic studies are described in *SI Appendix, Materials and Methods*.

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